PRODUCT: 47 bottles of *O. R. O.* at Lawton, Okla., together with a number of circulars entitled "O. R. O." Analysis showed that the product was a lime and sulfur solution.

LABEL, IN PART: (Bottle) "6 Ozs. or Over O. R. O. 14% light sulphur 2% hydrated lime 84% water (inert)."

NATURE OF CHARGE: Misbranding, Section 502 (a), certain statements on the label of the article and in the accompanying circulars were false and misleading. The statements represented and suggested that the article was effective in the treatment of unspecified diseases of poultry, running fits, mange, and other skin diseases of dogs, and coccidiosis and sorehead in chickens, turkeys, etc.; that it was effective to remove worms from poultry, hogs, dogs, or livestock of any kind; that it was effective to eliminate blue bugs and fleas on chickens and turkeys; and that it was effective in the treatment of poison ivy, prickly heat, ringworm, itchy scalp, scaly skin, chafed skin, rash, etc., of humans. The article was not effective for such purposes.

DISPOSITION: August 13, 1951. Default decree of condemnation and destruction.

DRUGS FOR VETERINARY USE *

3539. Misbranding of Dr. Jelen's Liquid Hog Medicine. U. S. v. 3 Pails, etc. (F. D. C. No. 31172. Sample No. 18916–L.)

LIBEL FILED: June 7, 1951. District of Minnesota.

ALLEGED SHIPMENT: On or about April 5, 1951, by Dr. Jelen's Veterinary Supply Corp., from Omaha, Nebr.

PRODUCT: 3 2-gallon pails and 5 1-gallon jugs of *Dr. Jelen's Liquid Hog Medicine* at New Ulm, Minn., together with a number of accompanying pamphlets entitled "Customer's Price List April 1950."

Examination indicated that the product consisted essentially of potassium arsenite, sodium hydroxide, sodium carbonate, sodium thiosulfate, sodium phosphate, potassium iodide (trace), creosote, anise oil, and licorice extract. Niacin (nicotinic acid) may have been present in the product but was not determined by analysis.

NATURE OF CHARGE: Misbranding, Section 502 (a), certain statements in the accompanying pamphlets were false and misleading. The statements represented and suggested that the article would aid in the treatment of necrotic enteritis ("necro"), and black scours; that it was helpful as a tonic in such cases; that it was a valuable aid in the feeding of poultry; that it would help one to keep his herd in better condition and free from "necro" at all times; and that it would help to keep the brood sow in good condition and to produce litters free from "necro." The article was not effective for the purposes stated and implied, and it was not capable of fulfilling the promises of benefit made for it.

DISPOSITION: July 27, 1951. Default decree of destruction.

3540. Misbranding of Dr. Mayfield poultry tablets. U. S. v. 23 Bottles, etc. (F. D. C. No. 29345. Sample Nos. 76130–K, 76131–K.)

LIBEL FILED: June 3, 1950, District of Minnesota.

ALLEGED SHIPMENT: On or about March 30 and April 12 and 27, 1950, by Dr. Mayfield Laboratories, Inc., from Charles City, Iowa.

*See also No. 3538.

PRODUCT: 23 1,000-tablet bottles and 39 500-tablet bottles of *Dr. Mayfield poultry tablets* at Sleepy Eye, Minn., together with a number of cards entitled "Poultry Disease Prevention Program" and a number of circulars entitled "Dr. Mayfield Poultry Tablets for Coccidiosis Control."

LABEL, IN PART: (Bottle) "Dr. Mayfield Poultry Tablets * * * Active Ingredients: Sodium Arsanilate (.35 grains arsenic per tablet expressed as metallic) Ammonium Phenolsulphonate and Boric Acid are therapeutically inactive."

NATURE OF CHARGE: Misbranding, Section 502 (a), certain statements on the bottle label and on the cards and circulars accompanying the article were false and misleading since the statements represented and suggested that the article was effective in the prevention and treatment of blackhead in turkeys and in the prevention of disease conditions of poultry when used as directed, whereas the article was not effective for the purposes stated and implied.

DISPOSITION: September 24, 1951. Dr. Mayfield Laboratories, Inc., claimant, having consented to the entry of a decree, judgment of condemnation was entered and the court ordered that the product be released under bond for relabeling under the supervision of the Federal Security Agency.

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PRODUCTS

SHIPPERS, MANUFACTURERS, AND DISTRIBUTORS

FEDERAL SECURITY AGENCY

FOOD AND DRUG ADMINISTRATION

NOTICES OF JUDGMENT UNDER THE FEDERAL FOOD, DRUG, AND COSMETIC ACT

[Given pursuant to section 705 of the Food, Drug, and Cosmetic Act]

3541–3549

DRUGS AND DEVICES

The cases reported herewith were instituted in the United States district courts by the United States attorneys, acting on reports submitted by the Federal Security Agency. This is a collection of cases adjudicated earlier than those now being recorded in current notices of judgment, but not published because complete records were not available immediately after the cases were terminated. Published by direction of the Federal Security Administrator.

CHARLES W. CRAWFORD, *Commissioner of Food and Drugs.*

WASHINGTON, D. C., *February 14, 1952.*

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